D2/D3 dopamine supports the precision of mental state inferences and self-relevance of joint social outcomes

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Abstract

Striatal dopamine is important to paranoid attributions, although its computational role in social inference remains elusive. We employed a simple game theoretic paradigm and a formal Bayesian model of intentional attributions to investigate dopamine D2/D3 antagonism on ongoing mental state inference following joint social outcomes. Haloperidol, compared to placebo, enhanced the impact of behaviour on beliefs about the harmful intent of a partner, and increased learning from recent encounters. These alterations caused significant changes to model covariation and induced negatively correlated self-interest and harmful intent attributions. Our findings suggest haloperidol improves flexibility in model-based beliefs about others and simultaneously reduces the self-relevance of observations. Our results may be representative of D2/D3 dopamine's role in supporting self-relevant mentalisation. Our data and model bridge theory between general and social accounts of value representation. We demonstrate initial evidence for the sensitivity of our model and short social paradigm to drug intervention and clinical dimensions, allowing distinctions between mechanisms that operate across traits and states.

Keywords

Dopamine; Belief; Mental State Inference; Paranoia; Mentalisation; Social Interaction

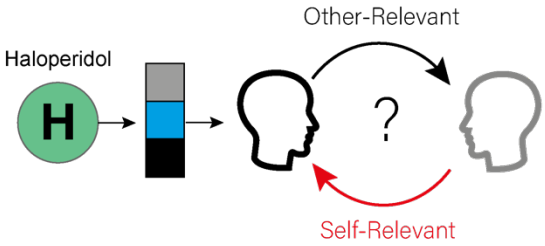
Data Availability

All data and code are available online:
https://github.com/josephmbarnby/Barnby_etal_2023_D2D3Modelling

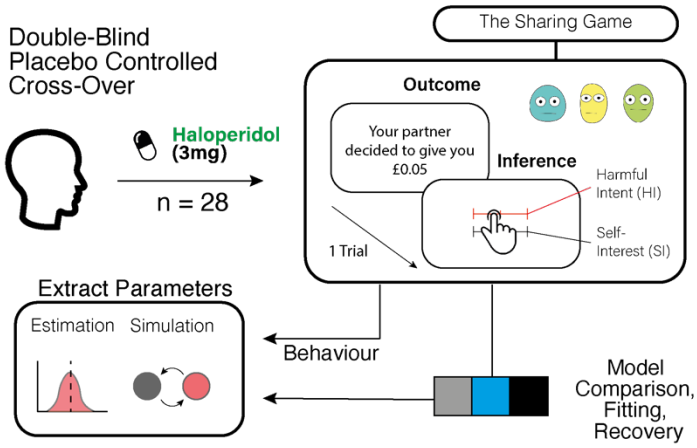
Graphical Abstract

Background

How can we better specify the causal role of D2/D3 dopamine on the computational mechanisms underlying mental state inferences in joint social contexts?

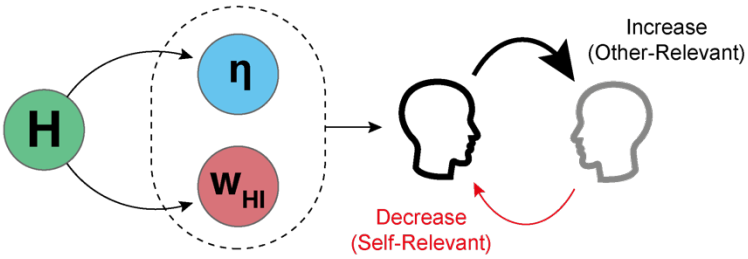


Methods & Analysis



Results

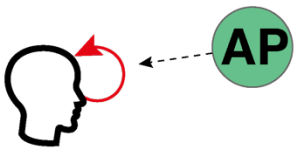
D2/D3 antagonism causally increases the impact of observations to change paranoid attributions, w_{HI}, and learning from experience, η.



Clinical Implications

D2/D3 Dopamine may support the self-relevance of information.

Anti**p**sychotics provide a break in recursive beliefs, reducing focus on the self following social outcomes.



Introduction

Dysregulated striatal dopamine has been identified as a key causal component in psychosis. Influential work proposed that striatal dopamine mediates aberrant salience leading to atypical perceptual experiences (Howes & Kapur, 2009; Kapur, 2004; Kapur et al., 2005) and more recent social-developmental models have highlighted the role of dopamine as a key point of convergence for a number of causal social and developmental factors, such as trauma, genetic vulnerability, and cannabis use (Howes & Murray, 2014). This has been supported by molecular and neuroimaging studies suggesting that developmental adversities (e.g., Dahoun et al., 2019; Egerton et al., 2016) increases pre-synaptic turnover of dopamine in striatal regions that may fuel the onset (Howes et al., 2011a; 2011b; Jauhar et al., 2019) and exacerbation (Laruelle et al., 1996; 1999) of psychosis symptoms.

Antipsychotics are the first-line treatment for psychosis and have good evidence for their efficacy (Schneider-Thoma et al., 2022). The exact mechanism by which their pharmacological effect reduces symptoms through the modulation of neurocognitive processes is still poorly understood. Understanding these pathways holds the promise of a better mechanistic understanding of psychosis and new targets for treatment development (Yip et al., 2022). A potential barrier to this process is that traditional verbally expressed mechanistic accounts of psychosis lack symptom-level and system-level specificity. Theorists must develop robust, falsifiable, empirical models of phenomena of interest (Haslbeck et al., 2022) and carefully map model parameters to the specific phenomenology of symptoms (Hitchcock et al., 2022) under a range of conditions.

Computational modelling bridges gaps between neuroscience and clinical expression (Huys et al., 2016; Montague et al., 2012), offering precise accounts of complex behaviour. Within this framework, dopaminergic alterations have been linked to core computational processes related to psychosis, such as aberrant salience (Howes et al., 2020), slower belief updating (Adams et al., 2013; Ashinoff et al., 2022), increased expectations of volatility (Diaconescu et al., 2020; Hauke et al., 2023; Reed et al., 2020), and model-based control (Mikus et al., 2022a).

One particularly disabling core symptom of psychosis is paranoia, the unfounded belief that others are trying to cause you harm (Freeman, 2016; Brakoulias & Starcevic, 2008). Psychologically, paranoia is characterised by sensitivities to interpersonal threat (Raihani & Bell, 2019), attributing negative outcomes to external, personal causes (Bentall et al., 1994), and overly complex mentalisation (Fonagy & Target, 1996). Developing computational theories to bridge the gap between the phenomenology and the neurocognitive mechanisms of paranoia requires particular considerations. Computational approaches in the social domain must sufficiently account for large, and often recursive, action spaces (Feldman Hall & Nassar, 2021; Velez & Gweon, 2021). These structural principles may be more appropriate for psychiatric symptoms which inherently involve alterations to interpersonal beliefs concerning the self and others (Barnby et al., 2023).

Models of intentional attributions – explicit inferences of the self about the mental state of others - allow for analyses that are theoretically related to ongoing paranoia. Current models include mechanistic explanations for perceived changes in the

harmful intent and self-interest that might motivate the actions of another. Prior work suggests high trait paranoia is associated with rigid priors about the harmful intent of partners, and a belief that a partner' actions are not consistent with their true intentions (Barnby et al., 2020; 2022a). Several predictions can be made concerning the influence of dopamine D2/D3 antagonism on paranoia. Synthetic, *in silico* models (Adams et al., 2022), neuroimaging evidence (Nour et al., 2018), prior predictions (Barnby et al., 2022a), and parallel psychopharmacological work (Mikus et al., 2022a; 2022b) predict that D2/D3 antagonism will increase belief flexibility and improve consistency of the self's model of others, which in turn should reduce self-relevant attributions of harmful intent following social outcomes. However, this has yet to be tested.

We examine the causal influence of D2/D3 dopamine receptor antagonism on computational mechanisms governing intentional attributions within a simple game theoretic context. Using a formal model of intentional attributions and an iterative Dictator game (Barnby et al., 2020a; 2022a), we test the impact of haloperidol, a D2/D3 antagonist, on paranoid beliefs using past data (Barnby et al., 2020b). We assess whether dopamine antagonism alters key computational processes involved in mental state inferences, allowing distinctions between trait representational changes (priors) and state learning processes (policy flexibility, uncertainty) along each dimension.

Methods

Participants

This study was approved by KCL ethics board (HR-16/17-0603). All data were collected between August 2018 and August 2019. Participants were recruited through adverts in the local area, adverts on social media, in addition to adverts circulated via internal emails.

Eighty-six participants were preliminarily phone screened. 35 participants were given a full medical screen. Thirty healthy males were recruited to take part in the full procedure. Two failed to complete all experimental days, leaving 28 participants for analysis. Inclusion criteria were that participants were healthy males, between the ages of 18 and 55. Participants were excluded if they had any evidence or history of clinically significant medical or psychiatric illness; if their use of prescription or non-prescription drugs was deemed unsuitable by the medical team; if they had any condition that may have inhibited drug absorption (e.g. gastrectomy), a history of harmful alcohol or drug use determined by clinical interview, use of tobacco or nicotine containing products in excess of the equivalent of five cigarettes per day, a positive urine drug screen, or were unwilling or unable to comply with the lifestyle guidelines. Participants were excluded who, in the opinion of the medical team and investigator, had any medical or psychological condition, or social circumstance, which would impair their ability to participate reliably in the study, or who may increase the risk to themselves or others by participating. Some of these criteria were determined through telephone check for non-sensitive information (age, gender, general understanding of the study, and overall health) before their full screening visit.

Procedure

This study was part of a larger study that assessed the role of dopaminergic modulation on personality, beliefs, and social interaction. Here, we focus on the role of dopamine antagonism and pre-synaptic increases in the attribution of mental state inferences during a Dictator game (described below; see Figure 1a).

The full procedure for participant screening is documented in a prior publication (Barnby et al., 2020). Briefly, participants who passed the brief phone screening were invited to attend an on-site screening day (see above). Participants were tested for drugs of abuse (SureScreen Diagnostics Ltd) and alcohol (breath test) prior to each experimental day and were excluded if any test was positive. Participants were given at least 7 days, but no more than two months, in between experimental days to allow for drug washout.

On experimental days, participants were randomised to be initially administered either a placebo or 3mg haloperidol in two capsules, and 10mg of domperidone in one capsule (3 caps total). After half an hour, participants were dosed a second time with either 150mg of co-beneldopa (herein referred to as L-DOPA) or placebo in two capsules. Participants would never receive haloperidol and L-DOPA in the same day.

The Sharing Game

Participants were asked to play a within-subjects, multi-trial modification on the Dictator game design used in previous studies to assess paranoia (Barnby et al., 2019; 2020), hereafter called 'The Sharing Game' (Figure 1b). In the game, participants played six trials against three different types of partner's who are assigned the role of Dictator. In each trial, participants were told that they have a total of £0.10 and their partner (the Dictator) had the choice to take half (£0.05) or all (£0.10) the money from the participant. Partner policies were one of three types: always take half of the money, have a 50:50 chance to take half or all of the money, or always take all of the money. These policies were labelled as fair, partially fair, and unfair, respectively. The order that participants were matched with partners was randomised. Each partner had a corresponding cartoon avatar with a neutral expression to support the notion that each of the six trials was with the same partner.

After each trial, participants were asked to rate on a scale of 1–100 (initialised at 50) to what degree they believed that their partner was motivated (a) by a desire to earn more (self-interest), and (b) by a desire to reduce their bonus in the trial (harmful intent). From the participants perspective, the actions of the partner can be framed as either arising from motivations that concern the gain of value for the partner irrespective of the participant (other-relevant) or arising from motivations that concern the loss of value for the participant (self-relevant).

After making all 36 attributions (two trial attributions for each of the six trials over three partners), participants were put in the role of the Dictator for six trials—whether to make a fair or unfair split of £0.10. Participants were first asked to choose an avatar from nine different cartoon faces before deciding on their six different splits. These Dictator decisions were not used for analysis but were collected to match

subsequent participants with decisions from real partners. Participants were paid a baseline payment for their completion, plus any bonus they won from the game.

Analysis

Behavioural data has been previously published (Barnby et al., 2020). Here, we apply three computational hypotheses which could explain the data, centred around a Bayesian model (Barnby et al., 2020; 2022) developed to explain mental state inference dynamics during social observation, where recursive, strategic social action is not a process of interest (Barnby et al., 2023). Model 1 allowed separate uncertainties and likelihood weights for each attribution, identical to our prior work (Barnby et al., 2020). In line with general theories of belief updating, Model 2 hypothesised that beliefs would be updating with the same likelihood weight. Model 3 hypothesised that prior beliefs share a single uncertainty free parameter over each distribution. Descriptions of the parameters within the winning model are in Table 1.

All computational models were fitted using the Hierarchical Bayesian Inference (HBI) algorithm which allows hierarchical parameter estimation while assuming random effects for group and individual model responsibility (Piray et al., 2019). This process is shown to be most robust to outliers versus non-hierarchical inference or standard hierarchical inference with fixed effects, and minimises parameter and model confusion (Piray et al., 2019). Parameters were estimated using the HBI in native space drawing from broad priors ($\mu_m = 0$, σ_m = 6.5; where $m = \{m_1, m_2, m_3\}$). This process was run independently for each drug condition due to the dependency of observations between conditions (the same participants were in each condition). Parameters were transformed into model-relevant space for analysis. All models and hierarchical fitting was implemented in Matlab (Version R2022B). All other analyses were conducted in R (version 4.2.3; x86_64 build) running on Mac OS (Ventura 13.0). All statistics are reported as: (X, 95%CI: Y, Z), where X is the regression coefficient, and Y and Z are the 95% lower and upper confidence intervals (CI), respectively. All dependent regressors were centred and scaled. To consider the uncertainty of estimates we conducted Bayesian paired sample t-tests to assess individual-level parameter changes. This used JAGS as a backend MCMC sampler (Bååth, 2014); differences in the mean are additionally reported with their corresponding effect sizes (Cohen's d) and posterior 95%HDI (High Density Interval). The raw output of this is listed in Table S1. Bayesian paired sample t-tests were also used to assess differences between attributional coupling over time. To note, in the original behavioural analysis (Barnby et al., 2020a) we excluded one extra participant due to their extreme trait psychometric paranoia score (leaving 27 participants), however trait paranoia was not the subject of this analysis, and hierarchical model fitting constrains group behaviour during parameter estimation. Nevertheless, for transparency, we include analytic estimates with the original 27 individual included for comparison. This did not change conclusions (Table S2).

We also sought to examine model covariance. Exploratory factor analysis used oblique rotation, including all parameter estimates for each individual within placebo and haloperidol conditions. Optimal factors were determined from observation of the scree plot and cross-validated model accuracy (Figure S9). Cross-validation used 10 folds with three repeats within a logistic general linear model. Parameter loadings and individual factor scores >|0.4| were retained for analysis.

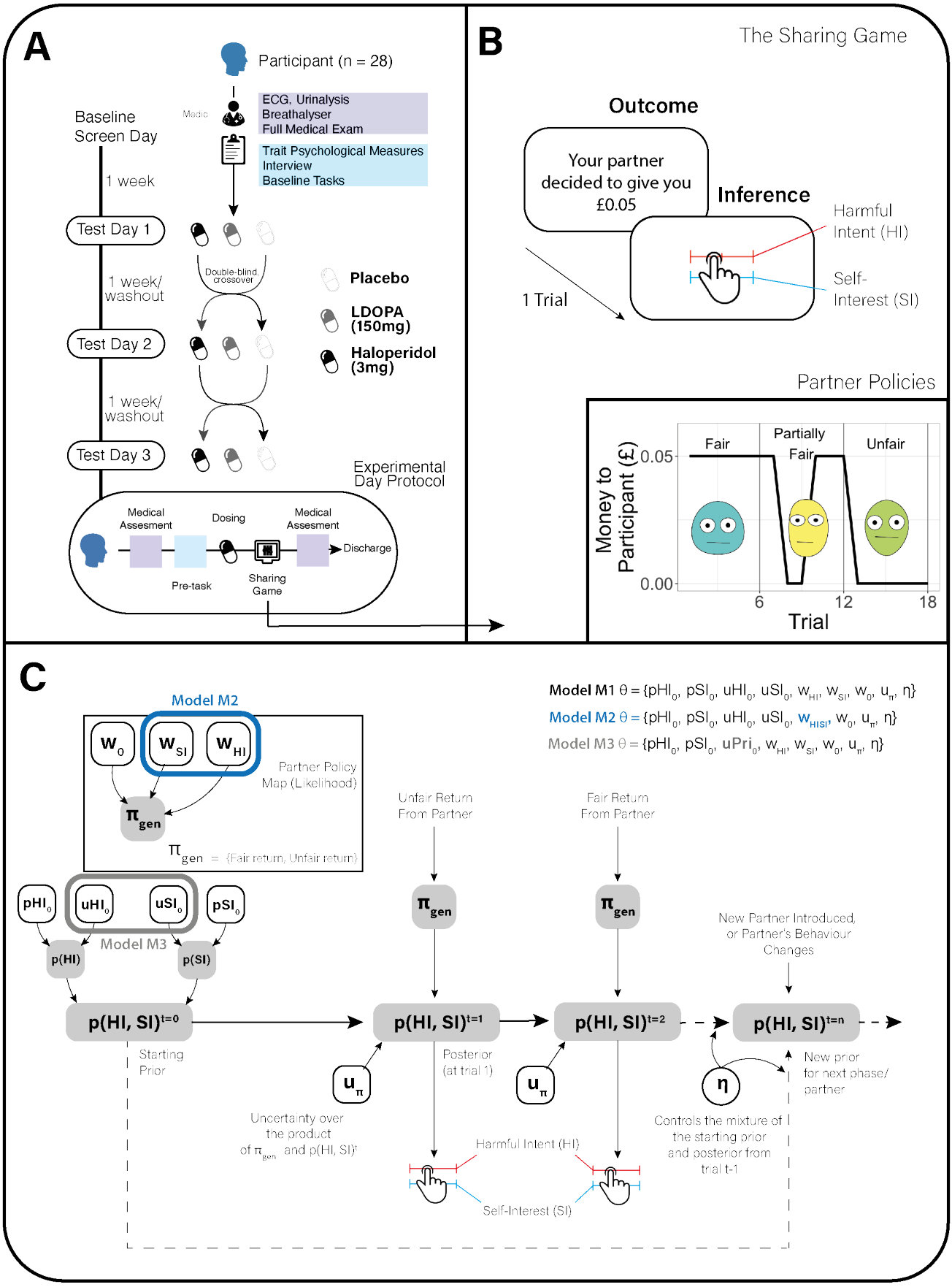

A
Participant (n = 28)
Medic
ECG, Urinalysis
Breathalyser
Full Medical Exam
Trait Psychological Measures
Interview
Baseline Tasks
Baseline Screen Day
1 week
Test Day 1
Double-blind, crossover
1 week/ washout
Test Day 2
1 week/ washout
Test Day 3
Placebo
LDOPA (150mg)
Haloperidol (3mg)
Experimental Day Protocol
Medical Assesment
Dosing
Medical Assesment
Discharge
Pre-task
Sharing Game
B
The Sharing Game
Outcome
Your partner decided to give you £0.05
Inference
Harmful Intent (HI)
Self-Interest (SI)
1 Trial
Partner Policies
Money to Participant (£)
0.05
0.00
Fair
Partially Fair
Unfair
6
12
18
Trial
C
Model M1 θ = {pHI0, pSI0, uHI0, uSI0, wHI, wSI, w0, uπ, η}
Model M2 θ = {pHI0, pSI0, uHI0, uSI0, wHISI, w0, uπ, η}
Model M3 θ = {pHI0, pSI0, uPri0, wHI, wSI, w0, uπ, η}
Model M2
w0
wSI
wHI
Partner Policy Map (Likelihood)
πgen
πgen = {Fair return, Unfair return}
pHI0
uHI0
uSI0
pSI0
Model M3
p(HI)
p(SI)
p(HI, SI)t=0
Starting Prior
Unfair Return From Partner
πgen
p(HI, SI)t=1
Posterior (at trial 1)
uπ
Uncertainty over the product of πgen and p(HI, SI)t
Harmful Intent (HI)
Self-Interest (SI)
Fair Return From Partner
πgen
p(HI, SI)t=2
uπ
New Partner Introduced, or Partner's Behaviour Changes
p(HI, SI)t=n
η
Controls the mixture of the starting prior and posterior from trial t-1
New prior for next phase/ partner

Figure 1. Experimental design and model space.

(A) Participants were entered into a double-blind, placebo-controlled, within-subject experimental design. (B) Participants engaged in a three-partner version of the sharing game. Here, partners were assigned the role of Dictator and on each trial could either take £0.10 for themselves (unfair outcome) or take £0.05 and give the participant £0.05 (fair outcome). Participant reported two types of attributional intent concerning the motivations of the partner after each outcome. These included harmful intent attributions and self-interest attributions. Partner order was randomised, and partner change was signalled. (C) Model space used to test whether dopamine manipulations were best explained by the full model (M1), a model that constrained policy updating to a single sensitivity parameter for each attribution (M2), or a model that constrained prior uncertainty to a single parameter (M3; Table 1).

Table 1. Winning model parameters and their causal structure. NB = number of discretised bins of each attributional response. Bin = binominal distribution with an added precision parameter, i.e. in the case of HI: $p(HI)^{t=0} \sim B(HI; \boldsymbol{pHI_0}, NB)^{1/\boldsymbol{uPri}}$

Parameter	Generative Purpose
$\boldsymbol{pHI_0}$	Magnitude of the prior that the actions of others are generally motivated by harmful intent (HI) toward the self, $p(HI)^{t=0}$. Increasing this parameter increases the belief that a partner is motived by harmful intent before any actions are observed.
$\boldsymbol{pSI_0}$	Magnitude of the prior that the actions of others are generally motivated by self-interest (SI) irrespective of the self, $p(SI)^{t=0}$. Increasing this parameter increases the belief that a partner is motived by self-interest before any actions are observed.
$\boldsymbol{uPri}$	Uncertainty over priors. Increasing this parameter broadens the prior distribution of both $p(HI)^{t=0}$ and $p(SI)^{t=0}$.
Prior	$p(HI)^{t=0} \sim Bin(HI; \boldsymbol{pHI_0}, \boldsymbol{uPri}, NB)$ $p(SI)^{t=0} \sim Bin(SI; \boldsymbol{pSI_0}, \boldsymbol{uPri}, NB)$ $p(HI, SI)^{t=0} = p(HI)^{t=0} p(SI)^{t=0}$ $NB = 9$
$\boldsymbol{w_0}$	Intercept of the likelihood matrix, π_{gen}, that calibrates the magnitude of attributional change when a fair or unfair action is made by a partner.
$\boldsymbol{w_{HI}}$	Impact of observation on beliefs that an outcome (r) is motivated by harmful intent. Increases in this parameter leads to greater influence of a partner's behaviour on attributions of harmful intent.
$\boldsymbol{w_{SI}}$	Impact of observation on beliefs that an outcome (r) is motivated by self-interest. Increases in this parameter leads to greater influence of a partner's behaviour on attributions of self-interest.
Likelihood	$\pi_{gen}(r = 0; HI, SI) = \sigma(\boldsymbol{w_0} + [\boldsymbol{w_{HI}} * HI - \delta] + [\boldsymbol{w_{SI}} * SI - \delta])$ $\pi_{gen}(r = 0.5; HI, SI) = 1 - \pi_{gen}(r = 0; HI, SI)$ $\delta = \frac{NB + 1}{2}$ $\sigma(x) = \frac{1}{1 + e^{-x}}$
Update	$p(\widehat{HI, SI})^t = \frac{\pi_{gen}(r; HI, SI) p(HI, SI)^{t-1}}{\sum_{HI', SI'} \pi_{gen}(r; HI', SI') p(HI', SI')^{t-1}}$
$\boldsymbol{u\pi}$	The consistency to which partners were believed to act in accordance with their character. Higher values reduce consistency, causing a partner's behaviour to have less impact on beliefs.
Consistency rule	$p(HI, SI)^t \propto p(\widehat{HI, SI})^{t \frac{1}{\boldsymbol{u\pi}}} + \xi$ $\xi = 0.02/NB^2$
$\boldsymbol{\eta}$	Controls the mixture of prior and posterior beliefs used as a starting point for each partner. Higher values indicate more reliance on information gathered from the last encounter, rather than reverting to prior beliefs. The product from the below equation, $\overline{p(HI, SI)}^{t=C}$ replaces $p(HI, SI)^{t-1}$ on the first trial with a new partner.
Change point	$\overline{p(HI, SI)}^{t=C} = p(HI, SI)^{t=0} * [1 - \boldsymbol{\eta}] + p(HI, SI)^{t=C} * \boldsymbol{\eta}$ $C = final\ action\ of\ an\ other$

Results

Behavioural results

Behavioural results were published previously (Barnby et al., 2020). To summarise, when averaged over all Dictators, haloperidol caused a reduction in harmful intent attributions versus placebo (-0.17, 95%CI: -0.28, -0.05), but L-DOPA did not. Haloperidol also increased self-interest attributions versus placebo (0.16, 95%CI: 0.05, 0.27), but L-DOPA did not. Unfair and partially fair Dictators both elicited higher harmful intent (Partially fair = 0.28, 95%CI: 0.16, 0.40; Unfair = 0.75, 95%CI: 0.63, 0.87) and self-interest attributions (Partially fair = 0.59, 95%CI: 0.63, 0.87; Unfair = 1.16, 95%CI: 1.05, 1.27) versus fair Dictators.

Model comparison and recovery

Bayesian hierarchical fitting and comparison identified that at the group level (Figure 2A), participants under placebo and haloperidol were best fitted by model 3. This model assumed agents use a single uncertainty over both attributional priors, although used separate likelihood weights to update their beliefs about their partners policy. In contrast, participants under L-DOPA were best fit by model 2. This model assumes participants hold individual uncertainties over their prior beliefs, although use the same likelihood weight to update both attributional dimensions.

For each condition we examined model generative performance and reliability. We extracted parameters for each individual under each condition according to the model that bore most responsibility for their behaviour (Figure 2B). We then simulated data for each participant with their individual-level parameters for each condition and model and re-estimated model comparison, recovered each model, generated attributions for each trial and dictator condition, and fitted regression models for main effects. Bayesian hierarchical fitting and comparison on simulated data demonstrated excellent similarity to group and individual level model responsibility and exceedance probabilities from real data (Figure S1A). Likewise, individual level parameters demonstrated excellent recovery (all Pearson *r* values > 0.71, p values ~ 0; Figure S1B, C & D). Simulated and real attributions demonstrated excellent recovery across all drug and dictator conditions (all Pearson *r* values > 0.62, p values ~ 0; Figure S1E). Simulated attributions also recovered the main effects of drug and dictator condition on attributional dynamics: haloperidol demonstrated reductions in harmful intent versus placebo (-0.26, 95%CI: -0.36, -0.16), but L-DOPA did not, and haloperidol increased self-interest attributions versus placebo (0.26, 95%CI: 0.15, 0.37), but L-DOPA did not.

We were most interested in examining the effect of haloperidol versus placebo in order to understand the mechanism behind the observed descriptive behavioural results. As model 3 achieved group-level dominance across both placebo and haloperidol conditions we were able to directly compare all individual-level, winning model parameters between-conditions $\{pHI_0, pSI_0, uPri, u\pi, \eta, w_0, w_{HI}, w_{SI}\}$ (Table 1; see below).

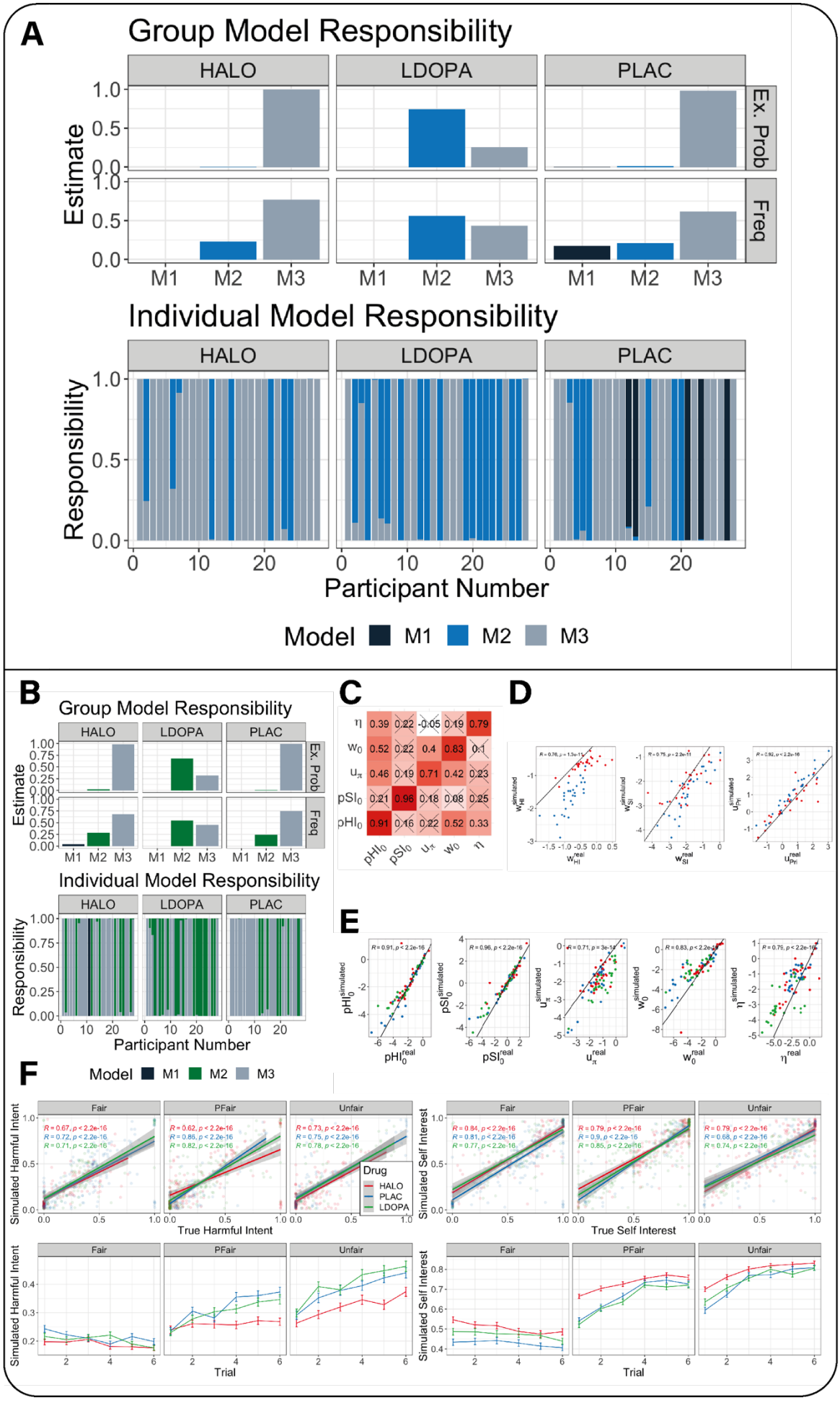

A
Group Model Responsibility
HALO
LDOPA
PLAC
Ex. Prob
Freq
Estimate
M1
M2
M3
Individual Model Responsibility
Responsibility
Participant Number
Model
B
C
D
E
F
Simulated Harmful Intent
True Harmful Intent
Simulated Self Interest
True Self Interest
Drug
Trial

Figure 2. Model comparison, recovery, and generative performance.

(A) Model responsibility across all three drug conditions. Greater model responsibility at the group and individual level indicates that a particular formulation was the most likely generative model to explain the data. Ex. Prob = Exceedance probability that a single model best defines group behaviour. Freq = Model frequency that each model is the best fitting model for participants. (B) Model recovery. All recovery analyses used n=28 synthetic participants – one for each real parameter set approximated from the data. The CBM algorithm correctly identified the correct model for most participants with trivial differences between model frequencies. (C) Correlation matrix of common parameters across all drug conditions for simulated (y axis) and real (x axis) data. All correlations were over 0.71 (p values < 0.001). 'X' indicates a non-significant association. (D) Individual correlations between common parameters across haloperidol and placebo conditions for simulated (y axis) and real (x axis) data. All correlations were over 0.71 (p values < 0.001). Black lines indicate the linear model of perfect association (r=1). (E) Individual correlations between common parameters across all drug conditions for simulated (y axis) and real (x axis) data. Black lines indicate the linear model of perfect association (r =1). (F) Top panel: Correlation between simulated and real harmful intent (right) and self-interest (left) attributions across all Dictator policies. Bottom panel: Simulated harmful intent (left) and self-interest (right) attributions for each drug condition and Dictator policy.

Haloperidol reduces the influence of priors and the precision of harmful intent

We examined the differences between individual level parameters within-subjects for haloperidol versus placebo (Figure 3A; see Figure S4 [Supplementary Materials] for effect sizes). This suggested that haloperidol increased reliance on learning about a partner just encountered, relative to pre-existing prior beliefs about partners in general (η; mean diff. = 0.15, 95%HDI: 0.03, 0.26; effect size = 0.66, 95%HDI: 0.22, 1.10). Haloperidol did not influence the consistency to which partners were believed to act in accordance with their character ($u\pi$).

Haloperidol increased learning flexibility over harmful intent attributions only. Haloperidol increased the impact of partner behaviour on harmful intent attributions (w_{HI}; mean diff. = 0.10, 95%HDI: 0.06, 0.13; effect size = 1.20, 95%HDI: 0.64, 1.75), but not over self-interest (w_{SI}); a partner's actions had more impact on a participant's beliefs about their true motivations of intentional harm. Haloperidol also caused the intercept of the policy matrix to be drawn toward 0, allowing greater updating parity for each partner action (w_0; mean diff. = 0.58, 95%HDI: 0.01, 1.10; effect size = 0.43, 95%HDI: 0.02, 0.82). The w_0 effect size should be treated with caution; the posterior distribution is within the region of practical equivalence (Figure S4).

We sought to further probe the model-based implications of drug differences on attributional flexibility in detail. Simulations on the marginal effect of w_{HI} on attributional dynamics are suggestive of its role in modulating the precision ($1/\sigma^2$) of attributions over all trials, irrespective of Dictator policy (Figure 3B). A regression model including w_{HI} as a linear term and w_0 as a quadratic term was most parsimonious compared to using w_0 as a linear term (AIC = 568 vs. 1123). There was a main effect of w_{HI} on the precision of harmful intent attributions (-6.13, 95%CI: -6.28, -5.97; effect size = -0.88, 95%CI: -0.92, -0.85). There was a small effect of w_0 within the same model (-0.06, 95%CI: -0.064, -0.056, effect size = -0.11, 95%CI: -0.14, -0.08). There was a significant but small interaction of w_0 and w_{HI} on the precision of harmful intent (-0.22, 95%CI: -0.25, -0.20; effect size = -0.05, -0.08, -0.02). Importantly, reductions in precision as a result of increased w_{HI} reduced harmful intent (-0.93, 95%CI: -0.95, -0.92; effect size = -0.13, 95%CI: -0.14, -0.13).

We also found evidence that a greater w_{HI} (cf. effect of haloperidol) may reduce precision most under conditions of ambiguity. Specifically, the precision of harmful intent attributions are lower in partially fair vs fair Dictators (-0.24, -0.33, -0.15; effect size = -0.24, 95%CI: -0.33, -0.15) and but unfair vs fair Dictators produced equivalent precision. Dictator policy interacts with w_{HI}: higher w_{HI} is associated with lower precision under partially fair vs. fair dictators (-0.77, 95%CI: -1.42, -0.42; effect size = -0.11, 95%CI: -0.21, -0.02). Thus, higher w_{HI} accentuates flexibility overall, but most in ambiguous social contexts where paranoia often flourishes. There was no interaction for unfair dictators vs. fair dictators (see Figure S5).

Haloperidol had no net significant influence on $pHI_0, uPri$, or pSI_0 (see Table S1). Individual parameter analysis suggests that haloperidol has a predominant net influence on the flexibility of belief updating about a specific context, here, that of our task. Under the influence of haloperidol, participants' assumptions about each new encounter are more amenable to change under the influence of recent encounters.

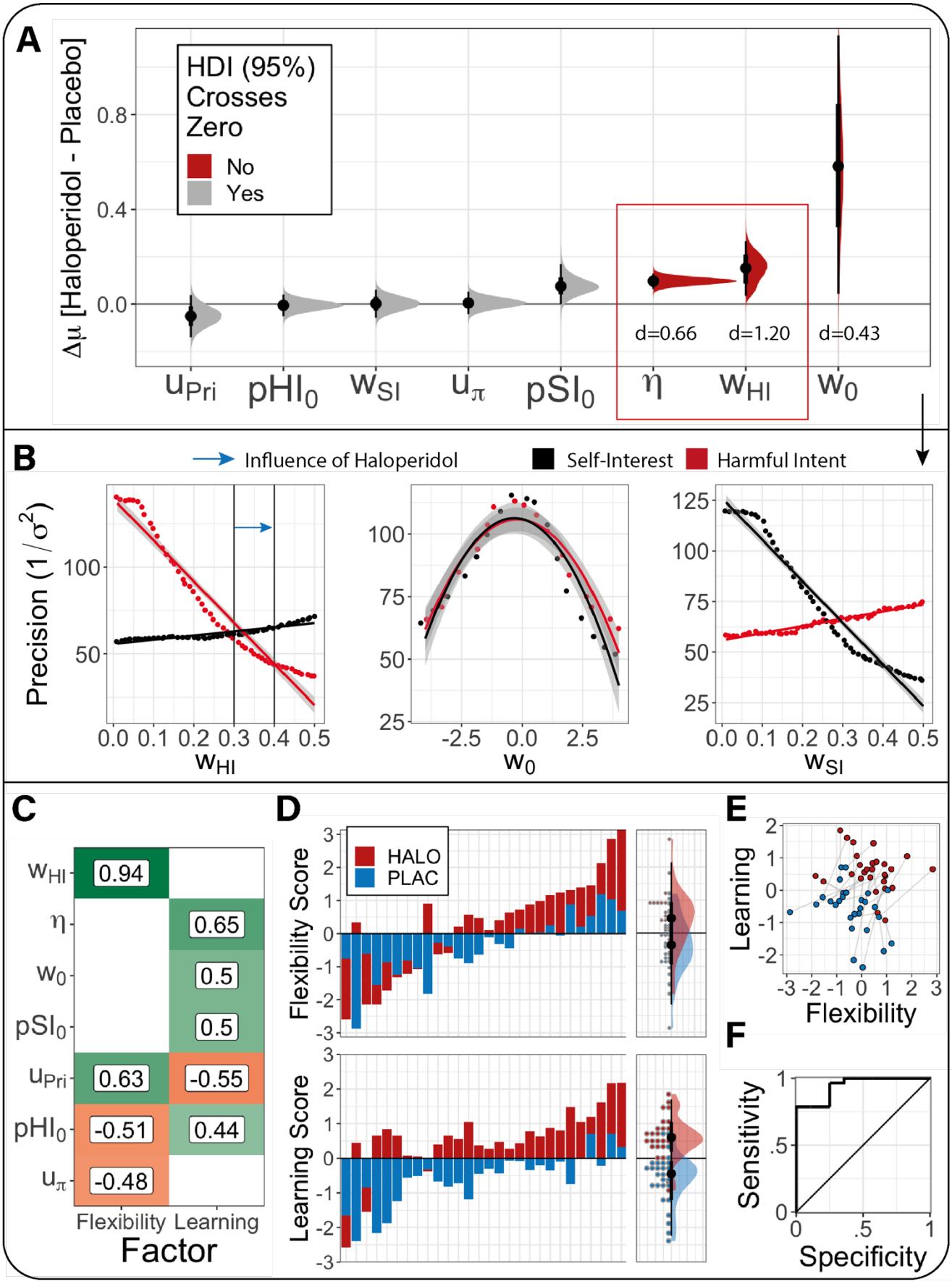

A
HDI (95%) Crosses Zero
No
Yes
Δμ [Haloperidol - Placebo]
0.8
0.4
0.0
d=0.66
d=1.20
d=0.43
uPri
pHI0
wSI
uπ
pSI0
η
wHI
w0
B
Influence of Haloperidol
Self-Interest
Harmful Intent
Precision (1/σ²)
100
50
0.0 0.1 0.2 0.3 0.4 0.5
wHI
100
75
50
25
-2.5 0.0 2.5
w0
125
100
75
50
25
0.0 0.1 0.2 0.3 0.4 0.5
wSI
C
wHI 0.94
η 0.65
w0 0.5
pSI0 0.5
uPri 0.63 -0.55
pHI0 -0.51 0.44
uπ -0.48
Flexibility Learning
Factor
D
HALO
PLAC
Flexibility Score
Learning Score
E
Learning
Flexibility
F
Sensitivity
Specificity

Figure 3. Influence of haloperidol on the winning model.

(A) Bayesian t-test results in assessing the difference and uncertainty of the change in mean ($\Delta\mu$) in parameter estimates between placebo and haloperidol. Red distributions indicate that the High-Density Interval (HDI) of the mean difference in distributions do not cross 0, suggesting reasonable certainty that the mean difference is not an artefact of statistical noise. 'd' values indicate the median effect size (Cohen's d) for each mean difference (See Figure S4 for distributions). The red box indicates parameters where effect size distributions were most robust, where the 95%HDI and lay outside of the region of probable equivalence with the null hypothesis. (B) Simulations of the marginal effect of likelihood parameters on the precision ($1/\sigma^2$) of harmful intent (red) and self-interest (black) attributions over all trial, controlling for Dictator style. Vertical lines are indicative of the median individual parameter estimates from both haloperidol and placebo groups, with the blue arrow indicating the difference from placebo to haloperidol. For trial-wise and within-Dictator precision changes see Figure S3; to note, simulations are consistent with the notion that w_{HI} reduces increases flexibility within and between contexts, accentuating smooth learning. To note, there was no significant correlation between w_0, w_{SI}, and w_{HI} in our parameter estimation from our real data (ps > 0.05; Figure S2) suggesting independent contributions of each to attributional dynamics. (C) Factor loading of each parameter on flexibility (factor 1) and learning (factor 2) dimensions. A loading filter of |0.4| was applied. Both of these factors were able to discriminate most effectively between drug conditions. w_{SI} is not featured in this plot as it was not meaningfully loaded onto either factor. (D) Factor scores for each individual participant (n=28) for both haloperidol (red) and placebo (blue) conditions ordered from low to high factor loading. The panels on the right of each graph demonstrate the marginal loading across participants. (E) Candyfloss plot of joint factor scores for each individual participant. Grey lines indicate that the same participant was responsible for each connected point under placebo (blue) and haloperidol (red) (F) Receiver Operating Characteristic curve describing the sensitivity and specificity of the combination of flexibility and learning factors on differentiating drug conditions. Area Under the Curve = 0.91. Sensitivity = 0.8. Specificity = 0.78.

Alterations to single parameters drive model covariation that differentiates haloperidol from placebo

From our analysis we can conclude that the model is accounting for the true observed data relatively well. Isolated parameter changes between conditions suggest this effect is primarily driven by increases in the impact of partner behaviour on beliefs about harmful intent, w_{HI}, and increased learning from experience, η. Considered separately, these key parameters did not fully explain how the model accounted for behaviour changes induced by haloperidol (Figure S4). We therefore sought to identify, through exploratory factor analysis, meaningful patterns over the covariation induced by Haloperidol.

We found that three factors best accounted for the data (Figure S9) with the first demonstrating the greatest eigenvalue (factor 1=2.82; factor 2=1.36; factor 3=1.13). K-fold cross-validation within a logistic model demonstrated that a two-factor solution provided the best median accuracy to discriminate between drug condition (mean accuracy = 0.86) and had the lowest AIC (40.3; see Fig S9). Each factor was able to predict drug condition independently (Factor 1 = 1.52, 95%CI: 0.50, 2.91; Factor 2 = 3.08, 95%CI: 1.72, 5.03), and there was a large effect found between conditions using Bayesian paired t-tests (factor 1: mean diff. = 0.76, 95%HDI = 0.37, 1.17; effect size = 0.94, 95%HDI = 0.35, 1.59; factor 2: mean diff. = 1.34, 95%HDI = 0.87, 1.85; effect size = 1.23, 95%HDI = 0.64, 1.84; Figure 3F).

Factor 1 (Flexibility; Figure 3C) was typified by high values of w_{HI}, and greater consistency between beliefs that a partner's actions are indicative of their true motivations, u_{π}. Factor 2 (Learning; Figure 3C) comprised high values of η, larger intercepts over the policy matrix, w_0, and higher values over priors pSI_0. pHI_0 and u_{Pri} were oppositely loaded onto each factor and would likely nullify each other in cases where participants scored strongly on both (Figure 3E). We note that pHI_0, and u_{Pri} load with slightly more absolute value on the Flexibility factor.

Haloperidol compresses the dimensionality of partner policies

Finally, we explored the impact of haloperidol on attributional coupling; this allows analysis into the dependency of different intentional components. To calculate this we estimated spearman correlations between harmful intent and self-interest for each trial across the sample, controlling for the type of Dictator policy affiliated. This revealed that while harmful intent and self-interest are attributed independently of one another under placebo (mean ρ[sd] = 0.03 [0.07]) replicating prior work (Barnby et al., 2020), under haloperidol they are negatively associated (mean ρ[sd] = -0.22 [0.08]), and this difference is significant (mean diff. = -0.26, 95%CI: -0.32, -0.20; effect size = 2.22, 95%HDI: 1.22, 3.24). This relationship was replicated using simulated model predictions (mean diff. = -0.25, 95%CI: -0.34, -0.17; effect size = -1.53, 95%HDI: -2.28, -0.78); see Figure 4A. There was evidence that the negative association induced under haloperidol decays over time (Pearson r = 0.52, p = 0.029). The same is not true under placebo (see Figure 4A). This interaction was not significant (regression coef. = -0.06, 95%CI: -0.12, 0.03). In sum, haloperidol causes harmful intent and self-interest attributions to become less independent. This means that those under haloperidol are more likely to believe someone must be more self-interested if they are perceived to be less intentionally harmful.

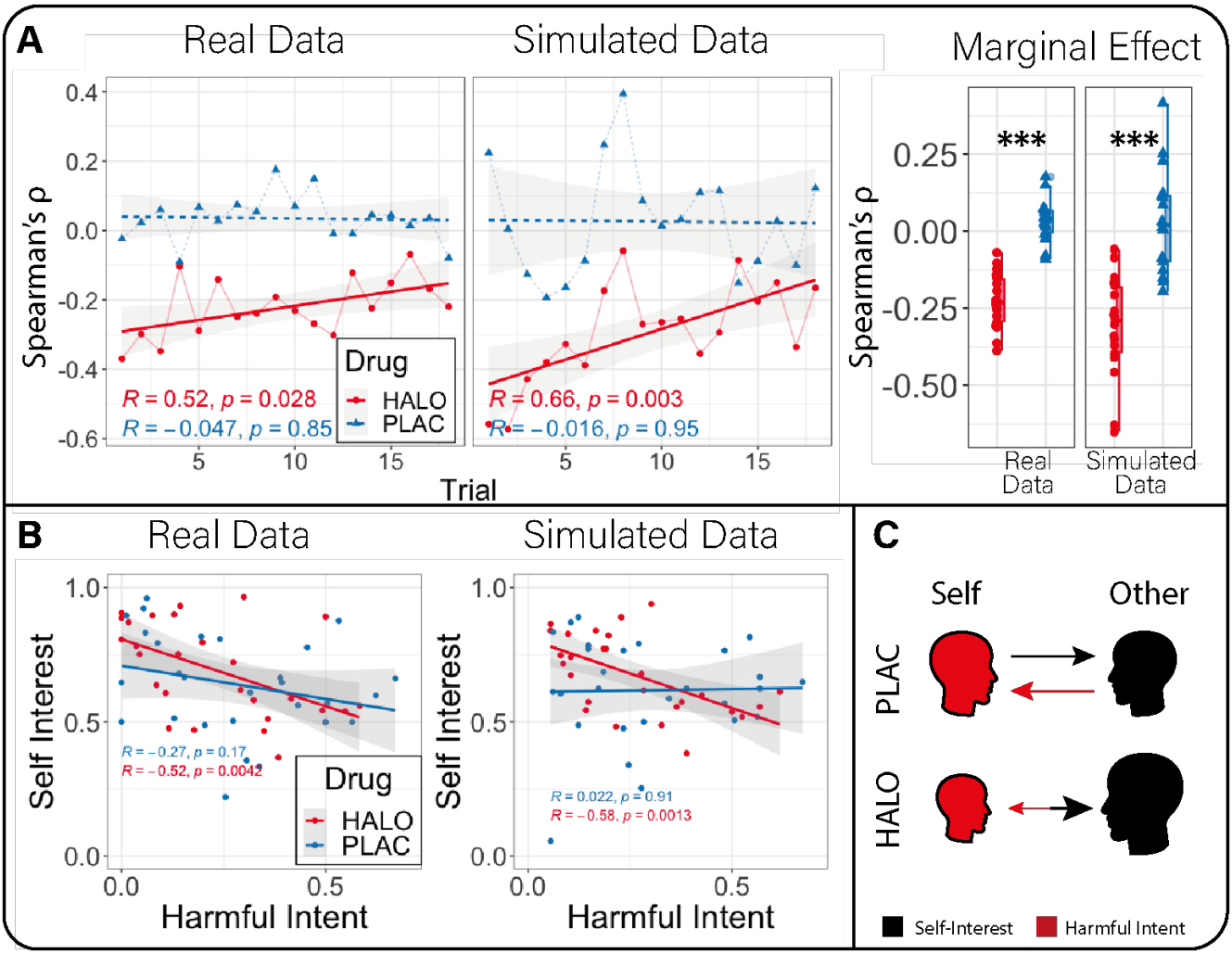


Figure 4. Association of mental state attributions between drug condition.

(A) In both real and simulated data, haloperidol (red) versus placebo (blue) induced a trial-wise negative association between harmful intent and self-interest which decayed over time. The right panel shows the marginal effect of trial-wise correlations between conditions. *** = p<0.001. (B) There was a general negative association between harmful intent and self-interest (Pearson correlation) found under haloperidol (red) for average attributions across all 18 trials. This was not true for placebo (blue). (C) Summary of main effects between drug conditions on self and other oriented intentional attributions following social outcomes. Both trial-wise and averaged associative analyses indicate that other-oriented attributions concerning self-interest of others (black) and self-oriented attributions concerning the harmful intent of others (red) are independent under placebo (PLAC) but coupled under haloperidol (HALO). Under haloperidol this coupling is biased toward exaggeration of other-oriented attributions and diminishment of self-oriented attributions.

Discussion

We sought to identify the computational mechanisms that explain how pharmacological alteration of dopamine function alters attributions of harmful intent, an important feature of paranoia, given our previous findings that haloperidol, but not L-DOPA, reduced harmful intent attributions and increased self-interest attributions in healthy participants (see Barnby et al., 2020a for previously published behavioural analysis). Here, we tested different computational hypotheses to account more mechanistically for these effects. The data were best fit by a model utilising a common uncertainty parameter over priors, but separate likelihood weights for updating attributions. Using this model, we found evidence that haloperidol reduced the precision of harmful intent (but not self-interest) attributions allowing more belief flexibility between partners. Haloperidol also increased the impact of learning from each encounter; participants relied less on their prior beliefs about the population as a whole. These individual parameter effects were embedded within covariational model alterations that together accounted for attributional change under haloperidol. These changes also caused negatively associated self-interest and harmful-intent attributions, suggesting a compression of attributions into a single interpersonal dimension under haloperidol. Together our findings indicate haloperidol promotes flexibility regarding attributions of harmful intent to others by reducing the perceived relevance of the actions of others to the self (Figure 5). In clinical environments this may allow space to reframe beliefs.

Our findings indicate a reduction in the influence of priors and more flexible beliefs under haloperidol. Previous research links tonic dopamine at D2/D3 receptors to efficient encoding of meaningful stimuli and Bayes optimality (Nour et al., 2018), cognitive control (Cools & D'Esposito, 2011), and sustained attention (Saeedi et al., 2016). Under the model-based, model-free control framework (Daw et al., 2005), recent work showed D2/D3 antagonism increased model-based control and decision flexibility (Mikus et al., 2022a) and increased belief flexibility during a trust game (Mikus et al., 2022b). This may be particularly useful in 'climbing out' of paranoia, where one is reluctant to take in positive information about others for fear of 'false reassurance'. At face value our results conform with previous work: under haloperidol, posteriors are more flexible and less influenced by priors, suggesting more confidence in beliefs about the motivation of partners. However, this general account does not explain why our data show asymmetric decreases in harmful intent and increases in self-interest.

One hypothesis is that haloperidol reduces the perceived self-relevance of outcomes under uncertainty. Social behaviours can rapidly develop complex action spaces. Humans try to reduce this uncertainty by relying on available heuristics, such as using self-preferences as an easily accessible prior belief about others (Anderson & Chen, 2002; Barnby et al., 2022b; Tarantola et al., 2017). When ambiguity increases, greater uncertainty about others (Barnby et al., 2020; 2022a; Hauke et al., 2023) and environments (Reed et al., 2022) can increase the perception of social threat. Our analysis suggests that haloperidol may attenuate the relationship between uncertainty and attributions of harmful intent by reducing the perceived self-relevance of others' actions; attributions of harmful intent, by definition, are inferences about the relevance of threat to the self from another. Given the role of the striatum and medial prefrontal cortex in regulating threat evaluation under stress

(Vaessen et al., 2015), this reduction in self-relevance may also interact with common neural implementations of self-other modelling (Nicolle et al., 2012); haloperidol may modulate the degree to which information is modelled as self- or other-relevant. The degree to which D2/D3 dopamine receptor function is specific to harmful intent or *all* attributions that are relevant to the self (e.g. altruistic intent of another) can be tested by including an extra dimension within our model; there are a number of hypotheses that can be made with such a modification (see Figure S7).

This pattern leads to a further, complimentary proposition: haloperidol may reduce self-relevance through reductions in the complexity or depth of recursive mentalising (how a self thinks about another's model of the self). In general, the ability to recursively mentalise is computationally expensive (Devaine et al., 2014; Guenounni & Speekenbrink, 2022; de Weerd et al., 2018); humans try to use cheaper, strategies when possible. Recursive mentalising is context dependent: in simple, competitive social scenarios humans are more likely to plan ahead more deeply and entertain recursive beliefs about another's model of the self (Goodie et al., 2012). Mentalisation gone awry has also been posited as a core driver of relationship difficulties in clinical populations: paranoia in borderline personality disorder and psychosis are explained as hyper-mentalisation – the inference of overly complex mental states based on sparse data (Fonagy & Bateman, 2006; Fonagy & Target, 1996; Sharp, 2014). An alteration in mechanisms that support self-relevant mentalising may explain our findings. This notion is consistent with reported amotivation under haloperidol, the role of D2/D3 receptors in promoting cognitive control (Cools & D'Esposito, 2011; Saeedi et al., 2016), and prior work on the causal role of D2/D3 antagonism in trust behaviours (Mikus et al., 2022b); reductions in the immediate value (and therefore relevance) for the self may facilitate longer term reciprocal trust behaviours without any need to engage deliberate reasoning about future outcomes. A core test of the hypothesis that D2/D3 dopamine is crucial for self-relevant, recursive mentalisation is to use models of hierarchical mentalisation in future experiments that allow estimation of recursive depth in joint social contexts.

The data presented here may be relevant beyond psychiatry. In behavioural economics, there have been several studies on the role of dopamine, reward, and decision making in both social and non-social contexts (Cox & Witten, 2019). Increasing dopamine availability has been shown to increase risky non-social decisions when self-gain is at stake (Rutledge et al., 2015), suggesting that dopamine may inflate the self-value of outcomes. Our data imply that this role of dopamine in modulating monetary value to the self may reflect a broader role in representing the self-relevance of stimuli. The direction of this relationship (self-relevance precedes self-value, or vice versa) is a fruitful target for future research. Our data may also be relevant to the role of dopamine in moral behaviour. In one study, boosting D2/D3 dopamine with pramipexole reduced generosity, especially with close others (Oroz Artigas et al., 2019). Our data compliments this work, suggesting that D2/D3 dopamine is involved in calibrating the valuation of self-gain in social decision-making.

On a theoretical level, our formal model distinguishes between computational changes that result from prior representational biases (e.g., higher trait paranoia) and acute state changes during social interaction where potential harm from others is a possibility (Figure 5). Previous modelling with the same task (Barnby et al.,

2020b) or a reversal variant of the task (Barnby et al., 2022) provided evidence that trait paranoia increases the magnitude of priors over harmful intent, the subsequent increase in the belief that the actions of others are not reflective of their true motivations, and a reduced willingness to believe that changes to a partner' behaviour are motivated by changes to their harmful intent. Naturally, this suggested that prior representations bias how social behaviour is interpreted. On the other hand, the present models suggest that haloperidol acts through increased reliance and impact of likelihoods on the formation of beliefs. Creating phenomenologically plausible formal models that are sensitive to different explanations of behavioural data has been a core aspiration of computational psychiatry (Hitchcock et al., 2022; Huys et al., 2016). Models like ours may be useful in distinguishing between longer term development and near-term alterations in learning that may explain paranoia. Model parameters are constant at the timescale of tasks while potentially evolving at the timescale of personal development, illness and recovery, while learning and inference can be dissected in the timescale of task conditions and trials. Much like prior work distinguishing interventions of representational change (psychotherapy) and emotion modulation (antidepressants; Nord et al., 2021) our model may support similar distinctions following intervention. We thus hypothesise that successful therapeutic use of haloperidol in paranoia will be associated with large changes in likelihood parameters described above but may leave intact, at least in the short term, prior beliefs about the harmful intent of others. D2/D3 independent processes may underpin ongoing vulnerability and may require further psychosocial learning.

We note some limitations. First, we did not use a patient population which means the extent to which the findings generalise to a population with persecutory delusions, rather than non-delusional paranoia, remains unclear. Second, we did not include any non-social comparator (e.g. model-based decision making or volatile environments) when assessing the role of haloperidol on cognition. This leaves a divide between how dopamine influences non-social cognition and mental state inferences. Prior work suggests some shared variance between more foundational computations (e.g. decision temperature, belief updating) and paranoia (Barnby et al., 2022; Nour et al., 2019; Reed et al., 2020). Replicating the present work with non-social comparators of our social task, e.g. using a slot machine partner, may help understand the relations between formal theories of general decision making and how this is expressed at a recursive and intentional level in the same individuals. Third, we did not use a design that probes how dopamine may facilitate generalisation of social knowledge outside of our game theory task. Prior work has demonstrated that representations about learned partners can pass on from one context to another (Guenounni & Speekenbrink, 2022); once a representation is learned using computationally intensive resources, a cheaper, heuristic model can be used. This related to the question of whether an associative model of updating may be more efficient once a policy is known, and given our findings, whether haloperidol causes a faster transition. Finally, despite the difference in model responsibility, we did not find any influence of L-DOPA on behaviour. This may be due to insufficient doses used to increase dopamine, or the unspecific activity of L-DOPA at specific sites.

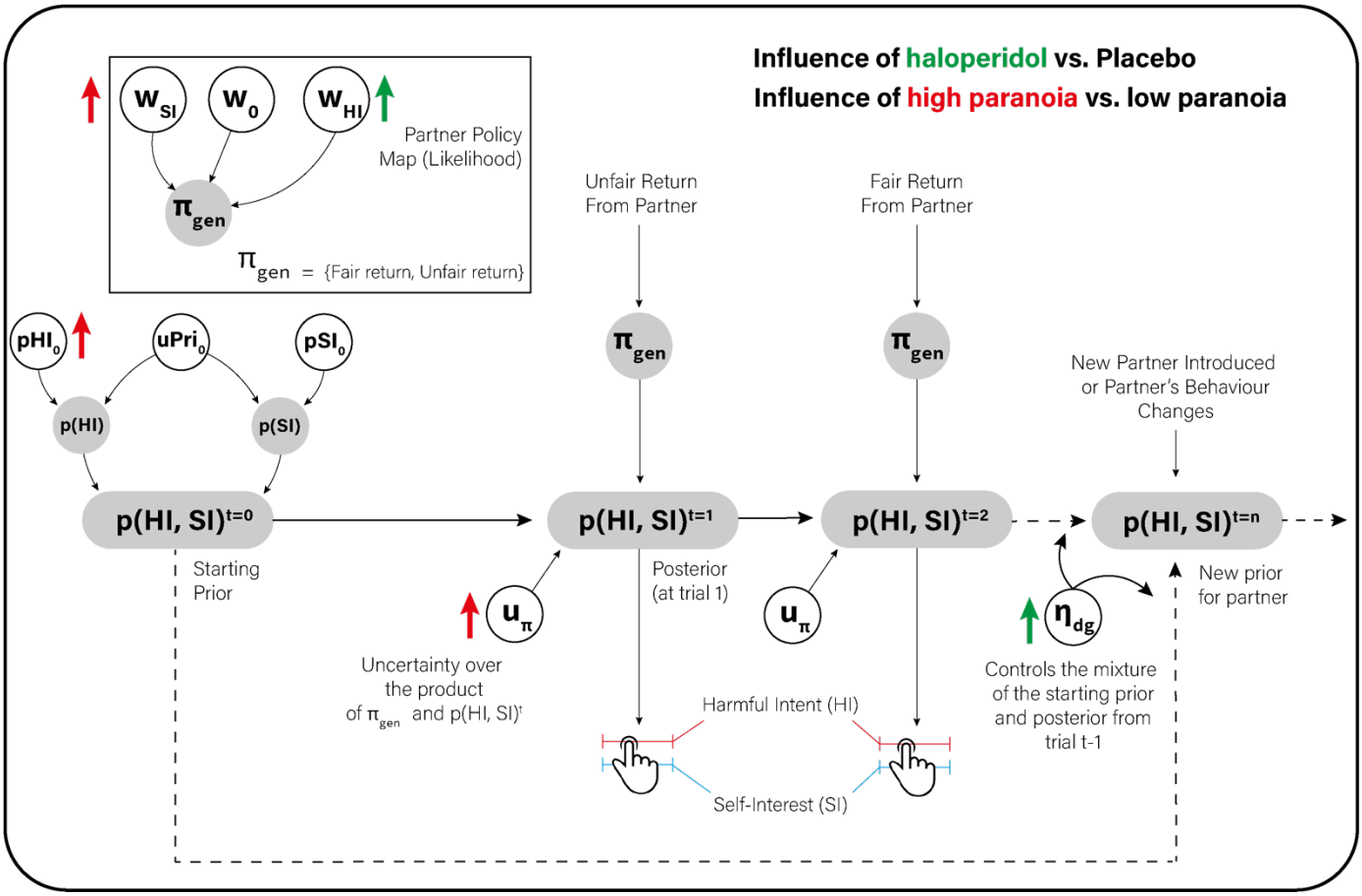


Figure 5. Summary of experimental parameter changes from current and past work.

(A) Experimentally observed effects on our model. Overall illustration of the impact of haloperidol on model parameters are illustrated in green. Prior results from the impact of high trait paranoia (Barnby et al., 2020; Barnby et al., 2022) are illustrated in red.

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